ACIDS + BASES

Electrolytes	- are	substances	that	dissolve	in water
\circ	and	form ions.			

Strong electrolytes - conduct electricity inagroom.

- ionic substances dissociate
- covalent molecular ionse:
- strong acids + bases,

Weak electrolytes - pool conductors

- small % of ionisation.
 - weak acids + bases.

Non electrolytes - no ions produced in water. - no current

Theores

Arrhenius Theory (1880)

- · acids provide H+ in aq.
- · bases provide OH in ag.
- · neutralisation Him + OHim > H20.(e)
- · restricted to aqueous solns

Brønsted - Lowry Theory (1923)

- · acids acts as proton (Ht) donors
- · bases act as proton acceptors.

H: OxH + Structure

small attach to polar substances

$$H_2O + NH_3 \rightarrow NH_4^+ + OH^-$$

Acia Base C.A CB.

When an acid donates a proton it leaves behind a conjugate base.

Strength

Strong acids completely ionise in water.

Weak acids partially.
Strong bases completely dissociate in water to on

Hydrolysis of Salts.

Basic anions - CH3000, HPQ, 2-, PO43-, SO4)-F , HCO3, CO3, C10, HS, CN-

(anion from weak acid)

Neutral anion: - CI, NO3, Br, I (anion from strong acid)

Acidic cations - NH4+, Al3+, Fe 3+ (cation from weak base netal ions.

Basic cattons - NONE (we need to know).

Neutral cution - group It II metals.

a) Nasoy

b) NH4CI



Hydrolysis of Water.

Self-ionisation of water.

Water weak conductor of electricity are to ionisation

OX

1 L pure water at 25°C contains 1.00×10⁻⁷ Ht. and 1.00×10⁻⁷ OH.

Water is neutral

As temp T so does Kw (pH changes)
but [OH-] = [H+] - neutral.

Example.

[H+] of a 3.50 mol L soln Bc(OH)2

[OH-] = 2 x [Ba(OH)] = 0.7 molL!

$$[H^{+}] = \frac{kw}{[OH^{-}]} = \frac{1 \times 10^{-14}}{0.700}$$

= 1.43×10-14 mol L-1

conc know

PH <7 acidic PH =7 neutral PH >7 basic.

Hydrolysis reactions of weak acids + bases

Acid Reaction

Weak acids - small equilibrium constant.

$$K_a = \frac{[H_3O^{\dagger}][CH_3COO^{\dagger}]}{[CH_3COOH^{\dagger}]}$$
acidity
constant

Base Reaction

Kb weak bases - small equilibrium constant pasicity contant

Polyprotic Acids + Bases.

Monoprotic acid - 1Ht/molecule of acid.

Diprotic acid - I ionisable Ht/molecule of acid.

e.g.
$$H_2SO_4 + H_2O \implies H_3O^+ + H_5O_4^-$$

 $H_3O_4^- + H_2O \implies H_3O^+ + SO_4^{2-}$

Triprotic acid - 3 ionisable 1+ / molecule of acid.

e.g.
$$H_3 PO_4 + H_2 O \ge H_3 O^4 + H_2 PO_4$$

$$H_2 PO_4^{-1} + H_2 O \ge H_3 O^4 + H_3 PO_4^{-2}$$

$$H_3 PO_4^{-1} + H_2 O \ge H_3 O^4 + PO_4^{-3}$$

Eg. Limiting reagent problem

Calculate the final pH of a soln by mixing 25.0 mL of 0.0750 moll HCI with 32.5 mL of 0.0675 moll kOH.

1. Write egm.

2. Calculate modes.

$$n(HCI) = c.V = 0.075 \times 6.027 = 1.875 \times 10^{-3} \text{ mol.}$$

 $n(KoH) = cV = 0.0675 \times 0.0327 = 2.194 \times 10^{-3} \text{ mol.}$

3. Ratio. 1:1

smallest amount HCI.

excess
$$n(koll) = n(koll) - n(lcl)$$
 newtralised.
= $2.194 \times 10^{-3} - 1.875 \times 10^{-3}$
= 3.188×10^{-4} mol.

4. Conc. of excess

$$C [KOIH] = \frac{n}{V}$$
= $\frac{3.188 \times 10^{-4}}{(6.0250 + 0.0327)} + \frac{sum of volumes}{volumes}$
= $5.5u \times 10^{-3} \text{ mol L}^{-1}$

5. pH calc.